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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.
09/237,291	01/25/99	YOUNG	J SYS-2068

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HM12/0410

EXAMINER

SCHMIDT, M

ART UNIT

PAPER NUMBER

1635

DATE MAILED: 04/10/00

Please find below and/or attached an Office communication concerning this application or proceeding.

Commissioner of Patents and Trademarks

Office Action Summary

Application No.

09/237,291

Applicant(s)

Young et al.

Examiner

Schmidt

Group Art Unit

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—The MAILING DATE of this communication appears on the cover sheet beneath the correspondence address—

Period for Response

A SHORTENED STATUTORY PERIOD FOR RESPONSE IS SET TO EXPIRE 3 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a response be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If the period for response specified above is less than thirty (30) days, a response within the statutory minimum of thirty (30) days will be considered timely.
- If NO period for response is specified above, such period shall, by default, expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to respond within the set or extended period for response will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133).

Status

- ☒ Responsive to communication(s) filed on 1/24/00.
- ☐ This action is FINAL.
- ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11; 453 O.G. 213.

Disposition of Claims

- ☒ Claim(s) 18-20, 23-27, 31-35, 37-44 and 46-51 is/are pending in the application.
- ☐ Of the above claim(s) _____ is/are withdrawn from consideration.
- ☐ Claim(s) _____ is/are allowed.
- ☒ Claim(s) 18-20, 23-27, 31-35, 37-44 and 46-51 is/are rejected.
- ☐ Claim(s) _____ is/are objected to.
- ☐ Claim(s) _____ are subject to restriction or election requirement.

Application Papers

- ☐ See the attached Notice of Draftsperson's Patent Drawing Review, PTO-948.
- ☐ The proposed drawing correction, filed on _____ is ☐ approved ☐ disapproved.
- ☐ The drawing(s) filed on _____ is/are objected to by the Examiner.
- ☐ The specification is objected to by the Examiner.
- ☐ The oath or declaration is objected to by the Examiner.

Priority under 35 U.S.C. § 119 (a)-(d)

- ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d).
- ☐ All ☐ Some* ☐ None of the CERTIFIED copies of the priority documents have been
- ☐ received.
- ☐ received in Application No. (Series Code/Serial Number) _____.
- ☐ received in this national stage application from the International Bureau (PCT Rule 1.7.2(a)).

*Certified copies not received: _____.

Attachment(s)

- ☒ Information Disclosure Statement(s), PTO-1449, Paper No(s). 7
- ☐ Interview Summary, PTO-413
- ☐ Notice of References Cited, PTO-892
- ☐ Notice of Informal Patent Application, PTO-152
- ☐ Notice of Draftsperson's Patent Drawing Review, PTO-948
- ☐ Other _____

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DETAILED ACTION

1. The use of the trademark RetroNectin has been noted in this application. It should be capitalized wherever it appears and be accompanied by the generic terminology.

Although the use of trademarks is permissible in patent applications, the proprietary nature of the marks should be respected and every effort made to prevent their use in any manner which might adversely affect their validity as trademarks.

Claim Rejections - 35 USC § 103

Claims 18-20, 23-27, 31-35, 37-44 and 46-51 are rejected under 35 U.S.C. 103(a) as being unpatentable over Murray et al. (US Patent 5,665,557), Nakahata (US Patent 5,861,315), Hoffman et al. (US Patent 5,744,361), Fei et al. (US Patent 5,635,387) or Davis et al. (US Patent 5,599,703), in view of Ku et al, Kobayashi et al, Ramsfjell et al (IDS Reference AK), Ohmizono et al, Szilvassy et al, Escary et al., or Bodine et al, and further in view of Tushinski et al (IDS Reference AN), Fletcher et al., Bello-Fernandez et al, Hatzfeld et al., and Hanenberg et al. (Nature Medicine Vol. 2, No. 8) or Henenberg et al. (IDS Reference AR).

The claims are drawn to methods of culturing human hematopoietic stem cells in media comprising mpl ligand, flt3 ligand (FL), c-kit ligand, IL3, TPO, IL6 and/or LIF, and further, genetically modifying said cells via contacting a retroviral, adenoviral or adeno-associated viral vector with the hematopoietic stem cells in culture. The claims were amended to further specify

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cultured with fibronectin, more specifically RiboNectin. Specifically, the claims are drawn to the following concentrations of growth factors (1) 0.1- about 500 ng/ml *mpl ligand* and *flt3 ligand* (claim 18), further, with 5- about 200ng/ml *c-kit ligand* (claim 19), and further, with 5- about 200 ng/ml *IL3* (claim 20); (2) 0.1- about 500 ng/ml *TPO*, *flt3 ligand* and *IL6* (claim 23 or claim 37), further with (a) 5- about 200 ng/ml of *LIF* (claim 24 or claim 39), or (b) 10 - about 100 ng/mL of *IL3* (claim 25 or claim 40), and further with 5- about 200 ng/mL of *c-kit ligand* (claim 27), or c) 5- about 200ng/mL of *c-kit ligand* (claim 26); (3) 5- about 200 ng/mL *TPO* and *FL* each, and about 10- about 100 ng/mL *IL6* (claim 31). The claims further specify the hematopoietic cells as CD34+Thy-1+ Lin- cells and the heterologous gene as a marker gene or a therapeutic gene. New claim 51 specifies the cells as human hematopoietic with CD34+Thy-1-.

Murray et al. (US Patent 5,665,557), Nakahata (US Patent 5,861,315), Hoffman et al. (US Patent 5,744,361), Fei et al. (US Patent 5,635,387) and Davis et al. (US Patent 5,599,703) are all relied upon to teach methods of isolating and culturing populations of human hematopoietic stem cells. They do not necessarily teach the combinations of cytokines taught in the instant specification for growth of the stem cells in culture nor do they necessarily teach retroviral mediated gene transfer into the isolated and cultured populations of hematopoietic stem cells. Specifically they teach for instance: (1) 10ng/ml IL-3, 2ng/mL GM-CSF, 100ng/mL SF and 2 units/mL of EPO (Murray et al., col. 5-6 and col. 16, lines 20-22); (2) 50ng/mL IL-6 and/or 100ng/mL SCF (Nakahata, col. 6, line 38); (3) 10-500ng/mL c-kit ligand (MCF) and at least one of IL-3 (.5-2ng/mL), GM-CSF (.1-1 ng/mL), IL-1 (1-10 U/mL), or IL-6 (.5-10 ng/mL) (Hoffman

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et al., col 4, lines 50-60); (4) 50 ng/mL SGF, 50 ng/mL IL#, 20 ng/mL GC-SF (Fei et al., col 18, lines 65-67 and col. 19, lines 1-23); and (5) 0.1-20 ng/mL GM-CSF, 1-200 ng/mL IL3, 5-500 ng/mL SCF and/or 1- 100 ng/mL IL6 (Davis et al., col 9, lines 11-32).

Ku et al., Kobayashi et al., Ramsfjell et al. and Ohmizono et al. are further relied upon to teach the effects of Tpo (mpl-ligand) and flt-ligand on human hematopoietic stem cells in culture. They do not necessarily teach retroviral mediated gene transfer in such cultured cell populations.

Szilvassy et al. and Escary et al. are further relied upon to teach the effects of LIF on hematopoietic stem cells in culture. They do not necessarily teach retroviral mediated gene transfer in such cultured cell populations.

Bodine et al. is further relied upon to teach the effects of SCF (c-kit ligand) and IL-6 on hematopoietic stem cells in culture. They do not necessarily teach retroviral mediated gene transfer in such cultured cell populations.

Tushinski et al., Fletcher et al., Bello-Fernandez et al. and Hatzfeld et al. are further relied upon to teach retroviral mediated transfer of genes into hematopoietic stem cells in culture. They do not necessarily teach the same culture condition instantly claimed.

Hanenberg et al. (Nature Medicine Vol. 2, No. 8) or Henenberg et al. (IDS Reference AR) are relied upon to teach the use of fibronectin (RetroNectin) to increase cell transformation of retroviral vectors.

It would have been prima facie obvious to one of ordinary skill in the art to culture human hematopoietic stem cells in mpl-ligand, flt3 ligand, c-kit-ligand, IL3, LIF, TPO, and/or IL6 since

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Murray et al. (US Patent 5,665,557), Nakahata (US Patent 5,861,315), Hoffman et al. (US Patent 5,744,361), Fei et al. (US Patent 5,635,387), Davis et al. (US Patent 5,599,703), Ku et al, Kobayashi et al, Ramsfjell et al, Ohmizono et al, Szilvassy et al, Escary et al., and Bodine et al. all teach methods for isolation and culturing of hematopoietic stem cells via addition of one or more of mpl-ligand, flt3 ligand, c-kit-ligand, IL3, LIF, TPO or IL6. It would have been further prima facie obvious to genetically modify said cultured cells via a retroviral vector since Tushinski et al., Fletcher et al., Bello-Fernandez et al. and Hatzfeld et al. all teach methods of retroviral mediated transfer into hematopoietic stem cells. Furthermore, it would have been prima facie obvious to one of ordinary skill in the art to use fibronectin to increase transduction efficiency of retrovirally transfected cells into hematopoietic cells as taught by Hanenberg et al. (Nature Medicine Vol. 2, No. 8) or Henenberg et al. (IDS Reference AR).

One of ordinary skill in the art would have been motivated to culture human hematopoietic stem cells via the methods taught by Murray et al. (US Patent 5,665,557), Nakahata (US Patent 5,861,315), Hoffman et al. (US Patent 5,744,361), Fei et al. (US Patent 5,635,387), Davis et al. (US Patent 5,599,703), Ku et al, Kobayashi et al, Ramsfjell et al, Ohmizono et al, Szilvassy et al, Escary et al., and Bodine et al. and further to transfect genes via retroviral vectors such as those taught by Tushinski et al., Fletcher et al., Bello-Fernandez et al. and Hatzfeld et al. and further using fibronectin (Retronectin) to increase transformation efficiency of the retrovirus into the hematopoietic cells as taught by Hanenberg et al. (Nature Medicine Vol. 2, No. 8) or Henenberg et al. (IDS Reference AR)

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One of ordinary skill in the art would have had an expectation of success to culture hematopoietic stem cells in mpl-ligand, flt3 ligand, c-kit-ligand, IL3, LIF, TPO, and/or IL6 for the effects taught by Murray et al. (US Patent 5,665,557), Nakahata (US Patent 5,861,315), Hoffman et al. (US Patent 5,744,361), Fei et al. (US Patent 5,635,387), Davis et al. (US Patent 5,599,703), Ku et al, Kobayashi et al, Ramsfjell et al, Ohmizono et al, Szilvassy et al, Escary et al., and Bodine et al. One of ordinary skill in the art would have had an expectation of success to genetically modify such human hematopoietic stem cells in culture via retrovirally mediated gene transfer as taught by Tushinski et al., Fletcher et al., Bello-Fernandez et al. and Hatzfeld et al. One of ordinary skill in the art would have further had an expectation of success to increase the transformation efficiency of retrovirus into human hematopoietic cells using fibronectin (RetroNectin) since it was known in the art that the recombinant peptide CH-296 (another name for fibronectin or RetroNectin) acts to bind retroviral vectors and cell surface proteins to increase efficient transfection (as taught by Hanenberg et al. (Nature Medicine Vol. 2, No. 8) or Henenberg et al. (IDS Reference AR) for instance).

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Any inquiry concerning this communication or earlier communications from the examiner should be directed to *Mary M. Schmidt*, whose telephone number is (703) 308-4471.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, *George Elliott, Ph.D.* may be reached at (703) 308-4003.

Any inquiry of a general nature or relating to the status of this application should be directed to the Group receptionist whose telephone number is (703) 308-0196.

M. M. Schmidt
April 9, 2000

A handwritten signature in cursive script, appearing to read "George C. Elliott".

George C. Elliott, Ph.D.
Supervisory Patent Examiner
Technology Center 1600